

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended): A human IL-18 substitution mutant, wherein said mutant comprises from ~~three~~ one to five amino acid substitutions in the sequence of SEQ ID NO:1, said substitutions being at from one to five amino acid residues chosen from the group consisting of: the cysteine at residue 38, the cysteine at residue 68, the cysteine at residue 76, the asparagine at residue 78, the glutamic acid at residue 121, the cysteine at residue 127, the leucine at residue 144, and the aspartic acid at residue 157.
2. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38 (SEQ ID NO:4).
3. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of asparagine at residue 78 (SEQ ID NO:5).
4. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of glutamic acid at residue 121 (SEQ ID NO:6).
5. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 3, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of leucine at residue 144 (SEQ ID NO:7).
6. (Currently amended): The human IL-18 substitution mutant ~~as claimed in Claim 1~~, wherein said mutant comprises the amino acid sequence of a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of aspartic acid at residue 157 (SEQ ID NO:8).
- 7.-8. Cancelled.

9. (Previously presented): A composition comprising a polypeptide conjugated to a water-soluble polymer, wherein the polypeptide is the human IL-18 substitution mutant set forth in SEQ ID NO:8.

10. (Previously presented): The composition as claimed in Claim 9, wherein the conjugation between the polypeptide and the polymer is covalent.

11. (Previously presented): The composition as claimed in Claim 9, wherein the water-soluble polymer is a member chosen from the group of: polyethylene glycol homopolymers, polyethylene glycol copolymers, polypropylene glycol homopolymers, poly(N-vinylpyrrolidone), poly(vinyl alcohol), poly(ethylene glycol-co-propylene glycol), poly(N-2-(hydroxypropyl)methacrylamide), poly(sialic acid), poly(N-acryloyl morpholine), and dextran.

12. (Previously presented): The composition as claimed in Claim 11, wherein the water-soluble polymer is unsubstituted.

13. (Previously presented): The composition as claimed in Claim 11, wherein the water-soluble polymer is substituted at one end with an alkyl group.

14. (Previously presented): The composition as claimed in Claim 11, wherein the water-soluble polymer is a polyethylene glycol homopolymer.

15. (Previously presented): The composition as claimed in Claim 14, wherein the polyethylene glycol homopolymer is monomethoxy-polyethylene glycol.

16. (Previously presented): The composition as claimed in Claim 15, wherein the monomethoxy-polyethylene glycol is chosen from the group of linear monomethoxy-polyethylene glycol and branched monomethoxy-polyethylene glycol.

17. (Previously presented): The composition as claimed in Claim 16, wherein the polyethylene glycol homopolymer has a molecular weight of from about 20,000 to about 40,000 daltons.

18. (Previously presented): The composition as claimed in Claim 17, wherein the polyethylene glycol homopolymer has a molecular weight of about 20,000 daltons.

19. (Previously presented): The composition as claimed in Claim 17, wherein the polyethylene glycol homopolymer has a molecular weight of about 30,000 daltons.
20. (Previously presented): The composition as claimed in Claim 17, wherein the polyethylene glycol homopolymer has a molecular weight of about 40,000 daltons.
21. Cancelled.
22. Cancelled.
23. (Withdrawn): A method of treating cancer in a patient by administering a therapeutically effective dose of the composition as claimed in Claim 18.
24. (Withdrawn): The method as claimed in Claim 23, wherein the cancer comprises an immunosensitive tumor chosen from the group of: renal cell carcinoma, myeloma, lymphoma, and melanoma.
25. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:4, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 38.
26. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:5, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 78.
27. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:6, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 121.
28. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:7, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 144.
29. (Previously presented): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:8, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 157.

30. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:9, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 144.

31. (Withdrawn): The composition as claimed in Claim 30, wherein the water-soluble polymer is chosen from the group of: linear polyethylene glycol homopolymer having a molecular weight of from about 20,000 to about 40,000 daltons and branched polyethylene glycol homopolymer having a molecular weight of from about 20,000 to about 40,000 daltons.

32. (Withdrawn): The composition as claimed in Claim 31, wherein the linear polyethylene glycol homopolymer has a molecular weight of about 20,000 daltons.

33. (Withdrawn): The composition as claimed in Claim 19, wherein the human IL-18 substitution mutant has the amino acid sequence set forth in SEQ ID NO:10, and wherein the mutant is conjugated to the water-soluble polymer at the cysteine at residue 157.

34. (Withdrawn): The composition as claimed in Claim 33, wherein the water-soluble polymer is linear polyethylene glycol homopolymer having a molecular weight of from about 20,000 to about 40,000 daltons.

35. (Withdrawn): The composition as claimed in Claim 34, wherein the linear polyethylene glycol homopolymer has a molecular weight of about 20,000 daltons.

36. Cancelled.

37. (Previously presented): A method of preparing a composition, said method comprising the steps of:

(a) obtaining the human IL-18 substitution mutant polypeptide of SEQ ID NO:8;
and

(b) conjugating the polypeptide with a functionalized water-soluble polymer at the cysteine at residue 157.

38. (Previously presented): The method of Claim 37, wherein the functionalized water soluble polymer is a member chosen from the group of: methoxy polyethylene glycol succinimidyl propionate, MW 20,000; methoxy polyethylene glycol succinimidyl propionate, MW 30,000; methoxy polyethylene glycol succinimidyl butanoate, MW 20,000; succinimidyl ester of

carboxymethylated methoxy polyethylene glycol, MW 20,000; methoxy polyethylene glycol aldehyde, MW 20,000; methoxy polyethylene glycol aldehyde, MW 30,000; methoxy polyethylene glycol hydrazide, MW 20,000; methoxy polyethylene glycol maleimide, MW 20,000; methoxy polyethylene glycol maleimide, MW 30,000; methoxy polyethylene glycol orthopyridyl disulfide, MW 20,000; methoxy polyethylene glycol orthopyridyl disulfide, MW 30,000; methoxy polyethylene glycol iodoacetamide, MW 20,000; and methoxy polyethylene glycol iodoacetamide, MW 30,000.

39. Cancelled.

40. Cancelled.

41. Cancelled.

42. (Previously presented): A method of improving the pharmacokinetics and pharmacodynamics of the human IL-18 substitution mutant of SEQ ID NO:8, said method comprising the step of conjugating the human IL-18 substitution mutant to a water-soluble polymer at the cysteine at residue 157.

43. (Withdrawn): The method as claimed in Claim 33, wherein the water-soluble polymer is a polyethylene glycol homopolymer.

44. Cancelled.

45. (Previously presented): The method as claimed in Claim 42, wherein the subcutaneous bioavailability is improved.

46. Cancelled.

47. (Previously presented): The method as claimed in Claim 42, wherein the subcutaneous bioavailability is improved, and binding to IL-18BP is reduced.

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2. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38 (SEQ ID NO:4).

3. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of asparagine at residue 78 (SEQ ID NO:5).

4. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 1, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of glutamic acid at residue 121 (SEQ ID NO:6).

5. (Withdrawn): The human IL-18 substitution mutant as claimed in Claim 3, wherein said mutant contains a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of leucine at residue 144 (SEQ ID NO:7).

6. (Currently amended): The human IL-18 substitution mutant ~~as claimed in Claim 1~~, wherein said mutant comprises the amino acid sequence of a serine in place of cysteine at residue 38, an aspartic acid in place of cysteine at residue 68, and a cysteine in place of aspartic acid at residue 157 (SEQ ID NO:8).

7.-8. Cancelled.